

PATENT COOPERATION TREATY

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

Commissioner
US Department of Commerce
United States Patent and Trademark
Office, PCT
2011 South Clark Place Room
CP2/5C24
Arlington, VA 22202
ETATS-UNIS D'AMERIQUE
in its capacity as elected Office

Date of mailing (day/month/year) 18 April 2001 (18.04.01)	Applicant's or agent's file reference Case #1887
International application No. PCT/US00/21872	Priority date (day/month/year) 10 August 1999 (10.08.99)
International filing date (day/month/year) 10 August 2000 (10.08.00)	Priority date (day/month/year) 10 August 1999 (10.08.99)
Applicant ASAOKA, Seiji et al	

1. The designated Office is hereby notified of its election made:

☒ in the demand filed with the International Preliminary Examining Authority on:

09 March 2001 (09.03.01)

☐ in a notice effecting later election filed with the International Bureau on:

2. The election ☒ was

☐ was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland Facsimile No.: (41-22) 740.14.35	Authorized officer Maria Kirchner Telephone No.: (41-22) 338.83.38
--	---

PATENT COOPERATION TREATY

PCT

REC'D 01 AUG 2001


WIPO

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

13

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference Case #1887		FOR FURTHER ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)	
International application No. PCT/US00/21872	International filing date (day/month/year) 10/08/2000	Priority date (day/month/year) 10/08/1999	
International Patent Classification (IPC) or national classification and IPC A61K7/06			
Applicant NATIONAL STARCH AND CHEMICAL INVESTMENT HOLDING CO			
<p>1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.</p> <p>2. This REPORT consists of a total of 5 sheets, including this cover sheet.</p> <p><input type="checkbox"/> This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).</p> <p>These annexes consist of a total of sheets.</p>			
<p>3. This report contains indications relating to the following items:</p> <ul style="list-style-type: none">I <input checked="" type="checkbox"/> Basis of the reportII <input type="checkbox"/> PriorityIII <input type="checkbox"/> Non-establishment of opinion with regard to novelty, inventive step and industrial applicabilityIV <input type="checkbox"/> Lack of unity of inventionV <input checked="" type="checkbox"/> Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statementVI <input type="checkbox"/> Certain documents citedVII <input checked="" type="checkbox"/> Certain defects in the international applicationVIII <input type="checkbox"/> Certain observations on the international application			
Date of submission of the demand 09/03/2001		Date of completion of this report 30.07.2001	
Name and mailing address of the international preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465		Authorized officer Renoth, H Telephone No. +49 89 2399 8589	



**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT**

International application No. PCT/US00/21872

I. Basis of the report

1. With regard to the **elements** of the international application (*Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17)*):

Description, pages:

1-77 as originally filed

Claims, No.:

1-9 as originally filed

2. With regard to the **language**, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.

These elements were available or furnished to this Authority in the following language: , which is:

- ☐ the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).
- ☐ the language of publication of the international application (under Rule 48.3(b)).
- ☐ the language of a translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3).

3. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in written form.
- ☐ filed together with the international application in computer readable form.
- ☐ furnished subsequently to this Authority in written form.
- ☐ furnished subsequently to this Authority in computer readable form.
- ☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.
- ☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. The amendments have resulted in the cancellation of:

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

5. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/US00/21872

(Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.)

6. Additional observations, if necessary:

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	1-9
	No:	Claims	
Inventive step (IS)	Yes:	Claims	1-9
	No:	Claims	
Industrial applicability (IA)	Yes:	Claims	1-9
	No:	Claims	

2. Citations and explanations see separate sheet

VII. Certain defects in the international application

The following defects in the form or contents of the international application have been noted:
see separate sheet

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/21872

Document WO 98/31329 is a patent family member and also mentioned on the front page of US 6,126,929, which is cited in the international search report.

The WO document has a publication date of 23.07.1998 and is, therefore, to be considered as a document of the state of the art according to Rule 33.1(a) PCT.

ad ITEM V

Reasoned statement under Article 35(2) with regard to novelty and inventive step.

Novelty, Article 33(2) PCT

The documents of the international search report disclose cosmetic compositions comprising polymers, e.g. polyurethanes, as film forming agents in combination with other ingredients.

However, a cosmetic composition as defined in independent claim 1 of the present application comprising the combination of a amphoteric urethane resin and a water-soluble resin is not disclosed in any of the documents cited in the international search report.

Independent claim 1 is, therefore, considered to be new over the cited prior art.

Dependent claims 2-9 disclose preferred embodiments of the claim to which they refer and, thus, they also meet the requirements of the PCT in respect of novelty.

Inventive step, Article 33(3) PCT

WO 98/31329 (D1) is considered to represent the closest prior art. It discloses that the combination of film-forming polymers (e.g. polyurethanes) and non-film-forming particles leads to formulations for coating hair, having good cosmetic properties (feel and ease of disentangling) and satisfactory mechanical properties for maintaining the hair.

The object of the present application is to provide cosmetics which include antithetic properties, i.e. touch and durability.

**INTERNATIONAL PRELIMINARY
EXAMINATION REPORT - SEPARATE SHEET**

International application No. PCT/US00/21872

It is not considered to be derivable from D1 that a combination of an amphoteric polyurethane and a water-soluble resin as defined in independent claim 1 solves this problem.

Independent claim 1 is, thus, considered to involve an inventive step.

Dependent claims 2-9 disclose preferred embodiments of the claim to which they refer and, thus, they also meet the requirements of the PCT in respect of inventive step.

ad ITEM VII

It is not possible to compare the examples and the comparative examples (cf. pages 29-75), since the disclosed tables do not contain any evaluation standard symbols (as well as the test explanations).

PCT

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference Case #1887	FOR FURTHER ACTION see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.	
International application No. PCT/US 00/ 21872	International filing date (day/month/year) 10/08/2000	(Earliest) Priority Date (day/month/year) 10/08/1999
Applicant NATIONAL STARCH AND CHEMICAL INVESTMENT HOLDING CO		

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 2 sheets.



It is also accompanied by a copy of each prior art document cited in this report.

1. Basis of the report

- a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.



the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

- b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :



contained in the international application in written form.



filed together with the international application in computer readable form.



furnished subsequently to this Authority in written form.



furnished subsequently to this Authority in computer readable form.



the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.



the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. ☐ **Certain claims were found unsearchable** (See Box I).

3. ☐ **Unity of invention is lacking** (see Box II).

4. With regard to the **title**,



the text is approved as submitted by the applicant.



the text has been established by this Authority to read as follows:

COSMETICS CONTAINING AMPHOTERIC POLYURETHANES

5. With regard to the **abstract**,



the text is approved as submitted by the applicant.



the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No.



as suggested by the applicant.



because the applicant failed to suggest a figure.



because this figure better characterizes the invention.



None of the figures.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/21872

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K7/06 A61K7/48 A61K7/11 A61K7/15 A61K7/043
 A61K7/02 A61K7/032

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, PAJ, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 775 483 A (OREAL) 28 May 1997 (1997-05-28) claims 1,6,8,18 ---	1-9
X	EP 0 793 957 A (OREAL) 10 September 1997 (1997-09-10) claims 1,6-11,20 ---	1-6,8,9
E	US 6 126 929 A (MOUGIN NATHALIE) 3 October 2000 (2000-10-03) claims 1,5-9,32-39 -----	1-5,8,9

☐

Further documents are listed in the continuation of box C.

☒

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

20 November 2000

Date of mailing of the international search report

27/11/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+31-70) 340-3016

Authorized officer

Beyss, E

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 00/21872

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP *0775483	A	28-05-1997	FR 2740336 A	30-04-1997
			FR 2740330 A	30-04-1997
			BR 9604455 A	23-06-1998
			CA 2188691 A	28-04-1997
			JP 9169615 A	30-06-1997
			JP 2000136117 A	16-05-2000
			US 5972354 A	26-10-1999

EP 0793957	A	10-09-1997	FR 2745494 A	05-09-1997
			AT 175109 T	15-01-1999
			BR 9707855 A	27-07-1999
			CA 2199053 A	04-09-1997
			CN 1212617 A	31-03-1999
			DE 69700082 D	11-02-1999
			DE 69700082 T	27-05-1999
			ES 2129998 T	16-06-1999
			WO 9732566 A	12-09-1997
			JP 11507396 T	29-06-1999
			PL 328600 A	01-02-1999
			US 6010686 A	04-01-2000

US 6126929	A	03-10-2000	CA 2249478 A	23-07-1998
			EP 0909157 A	21-04-1999
			WO 9831329 A	23-07-1998

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
15 February 2001 (15.02.2001)

PCT

(10) International Publication Number
WO 01/10397 A1

- (51) International Patent Classification⁷: **A61K 7/06**, 7/48, 7/11, 7/15, 7/043, 7/02, 7/032 (74) Agents: DEC, Ellen, T. et al.; National Starch and Chemical Company, P.O. Box 6500, Bridgewater, NJ 08807 (US).
- (21) International Application Number: PCT/US00/21872 (81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.
- (22) International Filing Date: 10 August 2000 (10.08.2000)
- (25) Filing Language: English
- (26) Publication Language: English
- (30) Priority Data:
11/226558 10 August 1999 (10.08.1999) JP (84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
- (71) Applicant (*for all designated States except US*): NATIONAL STARCH AND CHEMICAL INVESTMENT HOLDING CORPORATION [US/US]; P.O. Box 7663, Wilmington, DE 19803-7663 (US).
- (72) Inventors; and
- (75) Inventors/Applicants (*for US only*): ASAOKA, Seiji [JP/JP]; 18-26-702, Kamiyamate-cho, Suita-shi, Osaka 565-0841 (JP). KOYAMA, Katsuya [JP/JP]; 5-4-1-202, Senzo, Itami-shi, Hyogo 664-0898 (JP). TSUZUKI, Toshitaka [JP/JP]; 3-7-46, Senba-nishi, Minoo-shi, Osaka 562-0036 (JP). HASHIMOTO, Tomohiro [JP/JP]; 2-10-3, Nishi-tada, Kawanishi-shi, Hyogo 666-0138 (JP).

Published:

— With international search report.

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: COSMETICS CONTAINING AMPHOTERIC POLYURETHANES

(57) Abstract: There is provided cosmetics including antithetic physical properties such as touch and durability. Cosmetics contain amphoteric urethane resin having carboxyl group(s) and tertiary amino group(s) in one molecule and water-soluble resin.

WO 01/10397 A1

COSMETICS CONTAINING AMPHOTERIC POLYURETHANES

Detailed Description of the Invention

The present invention relates to cosmetics containing amphoteric urethane resin and water-soluble resin.

5 It is customary to use water-soluble resins such as nonionic polyvinyl pyrrolidone (PVP), cationic acrylic resin or cellulose, anionic acrylic resin or polyvinyl acetate, amphoteric acrylic resin or polyvinyl acetate etc. as a base resin of cosmetics such as hair fixative. The hair fixative using these water-soluble resins as base resin is
10 advantageous in that it has a high curl retention and excellent durability.

 However, in order to obtain hair fixative which is excellent in durability as above-mentioned, it is necessary to use a water-soluble resin which has a high glass-transition temperature (T_g), which has a
15 drawback in that its feel, which is important for cosmetics, is not good and its touch is very poor.

 The present invention has been made in view of such circumstance with the object of providing cosmetics which include antithetic physical properties, i.e. touch and durability.

20 In order to attain the said object, the cosmetics of the present invention have a structure that contains amphoteric resin having carboxyl group(s) and tertiary amino group(s) in one molecule and water-soluble resin.

 The inventors of the present invention have found that in the

hair fixative having as a base resin amphoteric urethane resin containing carboxyl group(s) and tertiary amino group(s) in one molecule, originally antithetic physical properties such as setting ability and touch, combing ability and resistance to flaking are compatible
5 with each other by the elasticity and toughness that the urethane resin possesses. Furthermore, they have also found that for neutral water, the said hair dressing is excellent in resistance to water due to the ionic bonding of the carboxyl group(s) and the tertiary amino group(s), and, for surface-active agent solutions such as shampoo in hair
10 washability, due to dissociation of the said ionic bonding, and at the same time, the said cationic tertiary amino group(s) exhibit(s) a good adherence resulting from its interaction with the surface of negatively charged hair, and they have filed a patent application for the cosmetic resin compositions containing the said amphoteric urethane resin as
15 the principal ingredient (Japanese Patent Application No.H10-27595). Thus, the amphoteric urethane resin has extremely excellent physical properties, but on the other hand, there lies a problem in durability, and they ,(the said inventors), have repeated studies for the purpose of improvement of the durability of amphoteric urethane resin. As a
20 consequence, they have discovered that if water-soluble resin and amphoteric urethane resin are used together, the problem of the durability which is a weak point of amphoteric urethane resin can be solved by the use of water-soluble resin, and the problem of the touch which is a weak point of water-soluble resin can be solved by the use

of amphoteric resin, thus resulting in the obtaining of a cosmetic rendered with antithetic physical properties such as touch and durability, and they have attained to the present invention.

Additionally the durability may be further improved by using
5 nonionic resin, anionic resin, cationic resin or amphoteric resin as said water-soluble resins.

The introduction into the structure of the said amphoteric urethane resin of the structural unit which may be derived from ethylene oxide as a nonionic hydrophilic constituent, may provide
10 sufficient hydrophilic nature, which ensures improvement of hair washability especially when it is used as hair cosmetic material.

Additionally, the introduction of polysiloxane bond(s) into the structure of the said amphoteric urethane resin may improve the feel which may be felt especially when it is used as hair cosmetic material.

15 EXAMPLES

Next, the Examples of the present invention will be described.

The cosmetic materials in accordance with the present invention may be obtained by using amphoteric urethane resin containing carboxyl group(s) and tertiary amino group(s) in one
20 molecule, and, water-soluble resin.

The cosmetics in accordance with the present invention may be used as hair cosmetics such as foam hair fixative, gel-like hair fixative, aerosol spray hair fixative, pump spray hair fixative, or cream-like hair fixative; skin care cosmetics such as shaving cream, skin care

lotion, or sun screen lotion; and make-up cosmetics such as nail polish, mascara, or foundation, and above all, it can be suitably used as hair cosmetics.

The said amphoteric urethane resin having carboxyl group(s) and tertiary amino group(s) in one molecule may be prepared by making a prepolymer having isocyanate group(s) by reacting the polyol compound (component A), polyisocyanate compound (component B), and compound C having active hydrogen(s) and carboxyl group(s) with each other in an excess of isocyanate group(s), and allowing the said prepolymer containing isocyanate group(s) to react with the compound (component D) having active hydrogen(s) and tertiary amino group(s). Alternatively, the said amphoteric urethane resin may be also prepared by making a prepolymer having isocyanate group(s) by reacting component C and component D in reverse order, namely by allowing the said compound A, compound B, and compound D to react with each other in an excess of isocyanate group(s), and then, allowing the prepolymer containing the isocyanate group(s) to react with the said specific compound C. Such processes make it possible to produce amphoteric urethane resin more easily and safely than conventional processes. In the production process as set forth, if the reaction between the compounds A and B is made concurrently with reaction between the specific compounds C and D, the carboxyl group(s) in compound C and the tertiary amino group(s) in compound D will form salts earlier, and it follows therefore that the

resultant is insoluble in the reaction system, so that it will not react with isocyanate compounds even if it has an OH group, whereby the intended amphoteric urethane resin can not be produced.

Referring to the said polyol compound (component A), it can
5 be any element that can be utilized in the production of common type polyurethane. General examples are polyester polyol, polyether polyol, polycarbonate polyol, polybutadiene polyol, polyisoprene polyol, polyolefin polyol, ester polyacrylate polyol, etc. These components may be used separately or in conjunction with more than
10 two kinds thereof. Above all, polyester polyol and polyether polyol may be adequately used.

Examples of the said polyester polyol are compounds originated from condensation polymerization of at least one kind of dicarboxylic acid such as succinic acid, glutaric acid, adipic acid,
15 sebacic acid, azelaic acid, maleic acid, fumaric acid, phthalic acid, terephthalic acid; and at least one kind of polyhydric alcohol such as ethylene glycol, propylene glycol, 1,4-butanediol, 1,3-butanediol, 1,6-hexane diol, neopentyl glycol, 1,8-octane diol, 1,10-decane diol, diethylene glycol, spiro glycol, trimethylolpropane; or those obtained by
20 the ring opening polymerization of lactones.

Referring to said polyether polyol, in addition to the polyhydric alcohol which is used in the synthesis of said polyester polyol, a compound resulted from ring opening polymerization of phenols such as bisphenol A, or primary amines or secondary amines with cyclic

ether such as ethylene oxide, propylene oxide, oxetane, tetrahydrofuran may be employed, and a compound resulted from ring opening polymerization of polyoxyethylenepolyol, polyoxypropylenepolyol, polyoxytetramethylenepolyol, and bisphenol A with at least one
5 of propyleneoxide and ethyleneoxide, etc. (in the case of copolymer, any of block copolymer and random copolymer may be available) can be cited by way of example.

Referring to polyisocyanate compound (component B), no specific element will be defined; organic diisocyanate compounds
10 such as aliphatic di-isocyanate compound, cycloaliphatic diisocyanate compound, perfume di-isocyanate compound can be cited by way of example, and these compounds can be used separately or in conjunction with more than two kinds thereof.

Examples of the said aliphatic diisocyanate compounds are
15 ethylenedi-isocyanate, 2,2,4-trimethylhexamethylene diisocyanate, and 1,6-hexamethylene diisocyanate, etc. Example of the said cycloaliphatic diisocyanate compounds are hydrogenated 4,4-diphenylmethane diisocyanate, 1,4-cyclohexane diisocyanate, methylcyclohexylene diisocyanate, isophorone diisocyanate, and
20 norbornane diisocyanate, etc. Examples of the said perfume diisocyanate compounds are 4,4-diphenylmethane di-isocyanate, xylene diisocyanate, toluene di-isocyanate, and naphthalene diisocyanate, etc. Of all these compounds, 1,6-hexamethylene diisocyanate, isophosphorone diisocyanate, norbornane diisocyanate,

etc. are preferable in that they are excellent in resistance to light and inexpensive.

Referring to the said compound (component C) having active hydrogen(s) and carboxyl group(s), any compound which includes at least one active hydrogen and at least one carboxyl group in its molecule suffices for the occasion. So, no specific compound will be defined. Examples are dimethylol propionic acid (DMPA), dimethylol butanoic acid, polycaprolactonediol containing carboxyl group(s), etc. They can be used separately or in conjunction with more than two kinds thereof.

Referring to the said compound (component D) having active hydrogen(s) and tertiary amino group(s), any compound which includes at least one active hydrogen and at least one tertiary amino group suffices for the occasion. So, no specific compound will be defined. Examples are N-alkyldialkanolamine compounds such as N-methyldiethanolamine, N-butyldiethanolamine, and dimethyl-aminoethanol, etc. They may be used separately or in conjunction with more than two kinds thereof.

When prepolymer containing isocyanate group(s) is produced using said components, chain extenders or molecular weight inhibitors can be used for regulation of characteristic features of amphoteric urethane resin as end product.

Referring to the said chain extender, no specific article will be defined. Examples of the article are low-molecular polyol and amines ,

etc. Examples of the said low-molecular polyol are glycols such as ethylene glycol, propylene glycol, 1,4-butanediol, diethylene glycol, 1,6 hexanediol, spiroglycol, cyclohexyl dimethanol, hydrogenated bisphenol A, neopentylglycol, bis (-hydroxyethoxy) benzene, and
5 xylylene glycol; and triol such as trimethylolpropane, and glycerine , etc.. Examples of said amines are ethylene diamine, propylene diamine, piperazine, hydrazine, isophorodiamine, methylene (bis-o-chloraniline), and polypropylene glycol containing amino groups at both ends.

10 Examples of the said molecular weight inhibitor are polypropylene glycol containing an amino group at one end and the like.

Solvent may be used in producing the said amphoteric urethane resin, as needed. For example, it is preferable to utilize
15 organic solvent which can dissolve both raw materials and polyurethane to be prepared. Examples of the said organic solvent are amides such as N-methylpyrrolidone, dimethylformamide, and dimethylacetamide; ketones such as acetone and methyl ethyl ketone; esters such as ethyl acetate; and cellosolve acetate, and cellosolve
20 ether, etc.

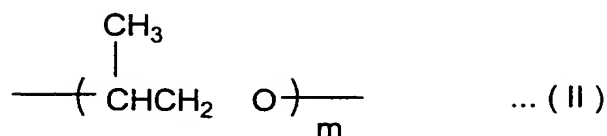
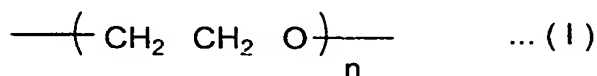
In the production of the said amphoteric urethane resin, the said resin can be provided with dispersibility in water by neutralizing the carboxyl group(s) or tertiary amino group(s) incorporated into the molecule by means of neutralizers, etc. Examples of neutralizers for

use in the said carboxyl group are triethylamine, trimethylamine, 2-amino-2-methyl-1-propanol, triethanolamine, potassium hydroxide, and sodium hydroxide, etc. Examples of neutralizers for the said tertiary amino group are acetic acid, hydrochloric acid, sulfuric acid, nitric acid, and dimethyl sulfate, etc.

Polymerizing catalysts known in the field of polyurethane may be used in the production of the said amphoteric urethane resin. Examples of the said catalyst that can be utilized are tertiary amine catalyst and organic metal catalyst, etc. Examples of the said tertiary amine catalyst are [2,2,2] diazabicyclooctane (DABCO), tetramethylenediamine, N-methylmorpholine, and diazacycloundecene (DBU), etc. Examples of the said organic metal catalyst are dibutyltin (tin) dilaurate, etc.

Referring to the said amphoteric urethane resin, it is preferable to use resins of the type having structural unit(s) which is/are derived from ethylene oxide (EO), in terms of hair washability.

Examples of the structural unit which is derived from the said EO are the EO unit as shown in the following general formula (I), and propylene oxide (afterwards referred to as PO) unit as shown in the following formula (II), and the EO unit is more preferable.



The said amphoteric urethane resin may have both EO and PO units. The ratio of the said EO unit to PO unit is preferably within the range of 10/0 to 2/8, and more preferably 10/0 to 4/6.

The repeated number n of the EO unit in the said general
5 formula (I) is set preferably within the range of 3 to 300, and more preferably 20 to 120. That is because if n is under 3, the EO unit(s) introduced into the amphoteric urethane resin is too little, so that the amphoteric urethane resin is not given sufficient hydrophilic nature, whereby there is a risk of the said resin obtaining insufficient hair
10 washability. Conversely, if n exceeds 300, the hydrophilic nature of the amphoteric urethane resin is too high, which may affect the moisture vapour resistance. Also, the repeated number m of the PO unit in the said general formula (II) is set preferably within the range of 3 to 300, and more preferably 20 to 120. If the said amphoteric urethane resin
15 has both the said EO unit and PO unit, $n + m$ preferably ranges from 3 to 300, and more preferably from 20 to 120.

The amphoteric urethane resin having the structural unit(s) which is/are derived from the said ethyleneoxide (EO) is produced by making a prepolymer having isocyanate group(s), for instance, by
20 allowing the polyol compound (component A), the polyisocyanate compound (component B), the polyethylene oxide derivative having active hydrogen(s), and a compound (component C) having active hydrogen(s) and carboxyl group(s) to react with each other in an excess of isocyanate group(s) and then, allowing the said prepolymer

containing isocyanate group(s) to react with a compound (component D) having active hydrogen(s) and tertiary amino group(s). Alternatively, this production process can be effective even by conducting the reaction with component C and the reaction with
5 component D in reverse order. In this case, the same compounds as aforementioned may be used as the said components A to D.

Referring to the polyethyleneoxide derivative having active hydrogen(s), any element that can introduce a structural unit that is derived from ethyleneoxide (EO) into the structure of the said
10 amphoteric urethane resin may be utilized. So, no specific element will be defined. Examples are polyoxyethylene glycol(PEG), and polyoxyethylene polyoxypropylene glycol (EOPO block copolymer), etc., and more preferably, preoxyethylene glycol. The said polyethylene oxide derivative of any of the following types can be
15 utilized; both-end OH group introduction type, both-end NH₂ group introduction type, one-end OH group introduction type, and one-end NH₂ introduction type. Amphoteric urethane resin having EO unit(s) in the main chain can be obtained by the use of the said both-end OH group introduction type or both-end NH₂ group introduction type.
20 Amphoteric urethane resin having EO unit(s) in the side chain(s) or end(s) can be obtained by the use of one-end OH group introduction type or one-side NH₂ group introduction type.

The molecular weight of the said specific polyethylene oxide derivative is preferably 200 to 20000, and more preferably 1000 to 10000.

It is preferable to use amphoteric urethane resin having
5 polysiloxane bond(s) in the structure thereof for much greater improvement of the feel.

Referring to the said polysiloxane bond(s), the repeated number n of siloxane bond (Si-O) is preferably within the range of 5 to 300, and more preferably 20 to 150. This is because if n is less than 5,
10 the proportion of siloxane bonds present in an amphoteric urethane resin being obtained may be too low, whereby sufficient effects which may be essentially acquired by the introduction of polysiloxane bond(s)s are hardly obtainable in touch, combing ability, etc. and conversely, with n exceeding 300, the compatibility with other
15 materials will deteriorate due to a high level of its hydrophobic nature, thus leading to poor reaction. Moreover, there is the possibility that the hydrophobic nature of a resultant polymer is at so a high level that it may hinder the adhesion to hair.

Amphoteric urethane resin having the said polysiloxane
20 bond(s) may be produced by making a prepolymer having isocyanate group(s), for example, by causing polyol compound (component A), polyisocyanate compound (component B), polysiloxane compound having active hydrogen(s), and a compound having active hydrogen(s) and carboxyl group(s) (component C) to react with each other in an

excess of isocyanate group(s), and then, reacting the resultant prepolymer containing isocyanate group(s) with a compound (compound D) having active hydrogen(s) and tertiary amino group(s). Alternatively, this production process can be effective even by
5 conducting the reaction with component C and the reaction with component D in reverse order. In this case, the same compounds as aforementioned may be used as said components A to D.

Referring to the polysiloxane compound having active hydrogen(s) that is used together with the said components A to D,
10 any compound that can introduce the polysiloxane bond(s) into the structure of the said amphoteric urethane resin suffices for the occasion. So, no specific compound will be defined. Examples of said compound are polydialkylsiloxanediol, polydialkylsiloxanemonool, polydialkylsiloxaneamine, and polydialkylsiloxanemonoamine, etc.
15 They may be used separately or in conjunction with more than two kinds thereof. For the alkyl group(s) which bond to Si of siloxane bond(s) of the said polydialkylsiloxanediol, etc., the number of carbons is preferably 1 to 10, and more preferably 1 to 5. It does not matter if the aforementioned polysiloxane compound incorporates the alkyl
20 group(s) which bond to Si of siloxane bond(s) and have a different number of carbons intermingled with each other. Specifically, examples of the said polydialkylsiloxanediol are polydimethylsiloxanediol, polymethylsiloxanediol, etc. Examples of the said polydialkylsiloxanemonool are polydimethylsiloxanemonool,

polymethylsiloxanemonool, etc. Examples of the said polydialkylsiloxanediamine are polydimethylsiloxanedimine, polymethylsiloxanediamine, etc. Examples of the said polydialkylsiloxanemonoamine are polydimethylsiloxanemonoamine, polymethylethylsiloxanemonoamine, etc.

Examples of the said polysiloxane compounds are those of both-end OH group introduction type, both-end NH₂ group introduction type, one-end OH group introduction type, and one-end NH₂ group introduction type, etc. The use of said compound of both-end OH group introduction type or both-end NH₂ group introduction type may bring about amphoteric urethane resin having polysiloxane bond(s) in the main chain, and the use of the compound of one-end OH group introduction type or one-end NH₂ group introduction type may bring about amphoteric urethane resin having polysiloxane bond(s) in the side chain(s) or at end(s).

In the cosmetics in accordance with the present invention, it is preferable to use amphoteric urethane resin as aqueous liquid. With the present invention, the aqueous liquid means the water dispersion in which amphoteric urethane resin is dispersed as well as the water solution in which amphoteric urethane resin is dissolved completely.

It is possible to make the water dispersion of said amphoteric urethane resin cross linkable by addition of a crosslinking agent such as a silane coupling agent. There is no restriction on adding various additives in order to impart storage stability to the said water

dispersion. Examples of the additives are protective colloid agent, anti-bacterial agent, mildewproofing agent, etc.

Referring to water-soluble resin which may be used together with the said amphoteric urethane resin, any of resins that can be
5 utilized as cosmetics suffices for this occasion. So, no specific resin will be defined. Every type of resin, i.e. nonionic, anionic, cationic, or amphoteric type can be employed. They may be used separately or in conjunction with more than two kinds thereof.

Examples of said nonionic resins are polyvinyl alcohol,
10 polyvinyl pyrrolidone [(Luviskol K-12,17,30,60,80,90 made by BASF Corporation), (PVP K-15,30,60,90,120 made by International Specialty Products)], vinylpyrrolidone-vinyl acetate copolymer [(Luviskol VA28,37,55,64,73 made by BASF Corporation), (PVP/VA-735,535,335,235,S-630 made by International Specialty Products),
15 (PVA-6450 made by OSAKA ORGANIC CHEMICAL IND.LTD), vinylpyrrolidone-vinyl acetate-vinyl propionate copolymer (Luviskol VA343 made by BASF Corporation), vinyl pyrrolidone-vinyl acetate-acrylamino acrylate copolymer, vinyl acetate/N-vinyl-5-methyl-2-oxazoline copolymer (Dowlex made by Dow Chemical Co.,Ltd),
20 hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropyl methyl cellulose, methyl cellulose, carboxymethyl cellulose, carboxyvinyl polymer, polyvinylformamide, and polyvinylacetamide , etc.

Examples of said anionic resins are vinyl acetate-crotonate copolymer [(Resin 28-1310 made by National Starch and Chemical

Co.), (Luviset CA made by BASF Corporation), crotonic acid-vinyl acetate-vinyl neodecanate copolymer (Resin 28-2930 made by National Starch and Chemical Co.), crotonic acid-vinyl acetate-vinyl propionate copolymer (Luviset CAP made by International Specialty Products), vinyl acetate-monobutyl maleate-isoboronyl acrylate copolymer (ADVANTAGE-CP made by International Specialty Products), N-octyl acrylamide/alkyl acrylates copolymer [(AMPHOMER HC made by National Starch and Chemical Co.), (Ultrahold 8, Ultrahold Strong made by BASF Corporation), vinyl pyrrolidone-acrylate-(meth) acrylic acid copolymer (Luviflex VBM35 made by BASF Corporation), acrylic resin alkanol amine liquid as (meth) acrylic acid-alkyl(meth)acrylate copolymer [(Pluscise series made by GOO Chemical Co., Ltd.), (Aniset KB-1000, KB-100H, NF-1000, HS-3000, AQ-2500 made by OSAKA ORGANIC CHEMICAL IND.LTD.), (Diahold made by Mitsubishi Chemical Corporation), alkyl acrylate-alkyl methacrylate-diaceton acrylamide-methacrylic acid copolymer (Pluscise L-53 made by GOO Chemical Co., Ltd.), methyl vinyl ether-maleic anhydride alkyl halfester copolymer [(Gantrez ES-225, ES-425, SP-215 made by International Specialty Products), (Anieres BEM-42S, VEM-22S made by OSAKA ORGANIC CHEMICAL IND.LTD.).

Examples of said cationic resins are (lower nitrogen) hydroxyethyl cellulose dimethyldiallylammonium chloride (Celquat L-200, H-100 made by National Starch and Chemical Co.), O- [2-

hydroxy-3-(trimethylammonio) propyl] hydroxyethyl cellulose chloride (Celquat SC240C, SC230M made by National Starch and Chemical Co.), vinyl pyrrolidone-quaternary dimethylaminoethyl methacrylate copolymer [(Gafquat 734, 755, 755N made by International Specialty Products), (H.C. Polymer 1S, 1N, 1NS, 1NP, 2, 2L, 3A, 5 made by OSAKA ORGANIC CHEMICAL IND.LTD.)], vinylimidazolium trichloride-vinylpyrrolidone copolymer (Luviquat FC370, FC550, FC905, HM552, MonoCP made by BASF Corporation), vinylpyrrolidone-dimethylaminoethyl methacrylate copolymer (Luviflex made by BASF Corporation), vinylpyrrolidone-alkylaminoacrylate-vinylcaprolactam copolymer (copolymer 845, 937, 958 made by International Specialty Products), vinylpyrrolidone-methacrylamidepropyl trimethylammonium chloride copolymer (Gafquat HS-100 made by International Specialty Products), alkylacrylamide-acrylate-alkylaminoalkylamide-polyethylene glycol methacrylate copolymer, polydimethylethylenepiperidinium chloride liquid (Marquat 100 made by CALGON Corporation), dimethyldiallylammonium chloride-acrylamide copolymer (Marquat 2200, 550 made by CALGON Corporation), and cationic guagum, etc.

Examples of the said amphoteric resin are octylamide acrylate-hydroxypropylacrylate-butylaminoethyl methacrylate copolymer (AMPHOMER 28-4910, LV-71 made by National Starch and Chemical Co.), N-methacryloyloxyethyl-N, N-dimethylammonium-N-methylcarboxybetaine/alkyl methacrylate copolymer [(Yukaformer

R205, R205S, SM, 301, 510, AM75, AM75S made by Mitsubishi Chemical Corporation), (RAM Resin-1000, 2000 made by OSAKA ORGANIC CHEMICAL IND.LTD.), acrylic acid-methacrylic acid-acrylic acid 2-hydroxypropyl-N, N-dimethylaminoethyl methacrylate-ethyl
5 methacrylate-diacetonacrylamide-vinylpyrrolidone copolymer and its alkanolamine liquid (AP Polymer 560 made by OSAKA ORGANIC CHEMICAL IND.LTD.), etc.

The average molecular weight of these water-soluble resins is preferably within the range of 100 to 10,000,000, and more
10 preferably from 10,000 to 5,000,000.

The blending proportion of the said amphoteric urethane resin to water-soluble resin in weight is preferably between 0.1/100 and 100/0.1, and more preferably 50/1 and 1/50.

In addition to the amphoteric urethane resins and water-
15 soluble resins, the cosmetics in accordance with the present invention may contain other ingredients which are commonly used in normal cosmetics, such as pigment, colouring matter, colouring material, perfume, surfactant, humectant, conservation agent, preservative, disinfectant, antioxidant, lubricant, thickening agent, ultraviolet
20 absorber, silicone polymer derivative, etc.

The cosmetics of the present invention may be produced in the following manner:

Production of hair cosmetics (foam hair fixative)

Aqueous liquid of the amphoteric urethane resins thus

obtained as mentioned above is blended with water-soluble resin; various kinds of surfactants such as polyoxy ethylene alkyl ether, coconut oil aliphatic acid diethanolamide, etc.; ethanol; de-ionized water, etc. in a predetermined proportion, and the resultant
5 compositions are mixed until they are made homogenous to produce a component X. Then, the addition of a component Y consisting of propellant (LPG), etc. to the said component X makes the intended foam hair fixative.

Production of hair cosmetics (aerosol spray-type hair fixative)

10 Aqueous liquid of the amphoteric urethane resins thus obtained as above-mentioned is combined with water-soluble resin, de-ionized water, dioctylsodium sulfosuccinate, ethanol, etc., in a predetermined proportion, and the resultant compositions are mixed until they are made homogenous to produce a component X. Then,
15 the addition of a component Y consisting of propellant (LPG), etc., to the said component X makes the intended aerosol spray-type hair fixative.

Production of hair cosmetics (gel-like hair fixative)

First, thickening agent, triethanolamine, ethanol, and de-
20 ionized water, etc. are combined with each other in a predetermined proportion, and the resultant compositions are mixed until viscous gel is formed into a component X. Then, aqueous liquid of the said amphoteric urethane resin is combined with water-soluble resin, ethanol, de-ionized water, etc. in a predetermined proportion to obtain

a component Y. Then, this component Y is added to said component X, and the resultant compositions are mixed until they are made homogeneous to produce the intended gel-like hair fixative.

Production of hair cosmetics (pumping spray-type hair fixative)

5 Aqueous water of the said amphoteric urethane resin is combined with water-soluble resin, dioctylsodium sulfosuccinate, ethanol, de-ionized water, etc. in a predetermined proportion, and the resultant compositions are mixed until they are made homogenous to produce the intended pumping spray-type hair fixative.

10 Skin care cosmetics such as shaving cream, skin care lotion, sunscreen lotion, etc.; and, make-up cosmetics such as nail polish, mascara and foundation, etc. can be produced in accordance with the general production processes for these cosmetics.

Next, the Examples will be described with reference to

15 Comparative Examples.

Prior to the description of the Examples and Comparative Examples, the following materials have been prepared.

Amphoteric urethane resin (a)

20 100g of isophorone diisocyanate (IPDI), 60g of polypropylene glycol (PPG, molecular weight: 1000), 5g of cyclohexyldimethanol (CHDM), and 38g of dimethylol butanoic acid(DMBA) were put into a glass flask with four openings and equipped with an agitator, a thermometer, a nitrogen duct, and a reflux condenser, then, 60g of ethyl acetate was added as a solvent, and the materials were heated

at a temperature of 80°C in the oil bath and allowed to react for four hours. Then, 2g of N-methyldiethanol amine and 30g of ethyl acetate were added, and the mixture was held for three hours for reaction purpose. Moreover, 30g of polypropylene glycol having one amino
5 group at one end thereof (Jeffamine M1000 made by HUNTSMAN CORPORATION) and 50g of ethyl acetate were added to the mixture, and the mixture was allowed to react for one hour, thereby obtaining a solution of polyurethane prepolymer with the residual NCO groups remaining therein. This polyurethane prepolymer with the residual
10 NCO groups was dispersed into 750g of water containing 16g of potassium hydroxide, and subjected to chain extension at 50°C for three hours for polymerization. Ethyl acetate was recovered from water dispersion thus obtained under a low pressure to obtain aqueous matters of amphoteric urethane resin without substantial
15 amounts of solvent.

Amphoteric urethane resin (b)

100g of isophorone diisocyanate (IPDI), 60g of propylene glycol (PPG, molecular weight:1000), 5g of cyclohexyl dimethanol (CHDM), 20g of polyoxyethylene glycol (PEG, molecular weight:1000),
20 and 36g of dimethylol butanoic acid (DMBA) were put into the glass flask with four openings and equipped with an agitator, a thermometer, a nitrogen duct and a reflux condenser, then, 60g of ethyl acetate as a solvent were added and the mixture was heated at a temperature 80 C in the oil bath to react for four hours. Then, 2g of N-

methyldiethanol amine and 30g of ethyl acetate were added, and the mixture was held for three hours for reaction purpose. Moreover, 30g of polypropylene glycol having one amino group at one end thereof (Jeffamine M1000 made by HUNTSMAN CORPORATION) and 50g of ethyl acetate were added to the mixture, and the mixture was allowed to react for one hour, thereby obtaining a solution of polyurethane prepolymer with residual NCO groups therein. This polyurethane polymer with residual NCO groups therein was dispersed in 750g of water containing 15g of potassium hydroxide and subjected to chain extension at a temperature of 50C for three hours for polymerization. Ethyl acetate was recovered from the resulting water dispersion under a low pressure to obtain aqueous matters of amphoteric urethane resin without substantial amounts of solvent but including ethyleneoxide chains in its structure.

15 Amphoteric urethane resin (c)

100g of isophorone diisocyanate (IPDI) and 3g of polydimethyl siloxanediol having two OH groups at one end thereof (molecular weight: 1000), were put into the glass flask with four openings and equipped with an agitator, a thermometer, a nitrogen duct and a reflux condenser and heated at a temperature of 80C in the oil bath to react for two hours. Then, 55g of polypropylene glycol (PPG, molecular weight: 1000), 10g of hydrogenated bisphenol A, and 36g of dimethylol butanoic acid (DMBA) were added together with 60g of ethyl acetate as a solvent, and the mixture was heated at 80C in

the oil bath to react for four hours. Then, 2g of N-methyldiethanol amine and 30g of ethyl acetate were added, and the mixture was allowed to react for another three hours. Moreover, 30g of polypropylene glycol having one amino group at one end thereof
5 (Jeffamine M1000 made by HUNTSMAN CORPORATION) and 50g of ethyl acetate were added to the mixture, and the mixture was allowed to react for another one hour, thereby obtaining a solution of polyurethane prepolymer with residual NCO groups therein. This polyurethane prepolymer with residual NCO groups therein was
10 dispersed in 750 g water containing 15g of potassium hydroxide, and subjected to chain extension at a temperature of 50C for three hours for polymerization. Ethyl acetate was recovered from the resulting water dispersion under a low pressure to obtain aqueous matters of amphoteric urethane resin without substantial amounts of
15 solvent but incorporating dimethyl siloxane chains in its structure.

Water-soluble resin (1) (nonionic)

Polyvinyl formamide.

Water-soluble resin (2) (nonionic)

Polyvinyl pyrrolidone (Luviskol K-90 made by BASF
20 Corporation).

Water-soluble resin (3) (anionic)

Crotonic acid-vinyl acetate-vinyl neodecanoic acid copolymer
(Resin 28-2930 made by National Starch and Chemical Co.)

Water-soluble resin (4) (anionic)

Acrylic resin alkanolamine liquid (Pluscise L-9909B made by GOO Chemical Co., Ltd.)

Water-soluble resin (5) (cationic)

- 5 Hydroxy ethylcellulose-dimethyl diallyl ammoniumchloride
(Celquat L-200 made by National Starch and Chemical Co.)

Water-soluble resin (6) (cationic)

- Vinyl pyrrolidone-N, N-dimethyl aminoethyl methacrylic acid
copolymer diethylsulfate (Gafquat 755N made by International
10 Specialty Products)

Water-soluble resin (7) (amphoteric)

Octylamide acrylate-hydroxypropyl acrylate-butylamino ethyl
methacrylate copolymer (AMPHOMER 28-4910 made by National
Starch and Chemical Co.)

- 15 Water-soluble resin (8) (amphoteric)

N-methacryl oyloxyethyl-N, N-dimethylammonium--N-
methyl carboxybetaine-alkyl methacrylate copolymer (Yukaformer SM
made by Mitsubishi Chemical Corporation)

Polyoxyethylene stearyl ether

- 20 NIKKOL BS-20 made by NIKKO CHEMICALS CO., LTD.

Coconut oil aliphatic acid diethanolamide

Amicor CDE-1 made by MIYOSHI OIL & FAT CO., LTD.

Diocetyl sodium sulfosuccinate

Monawet MO-70E made by MONA INDUSTRIES INC.

Thickening agent

Alkyl acrylate-polyoxy ethylene stearyl ether itaconate copolymer (Structure 2001 made by National Starch and Chemical Co.)

5 Hair cosmetics (foam hair fixative)Examples 1a to 24a, Comparative Examples 1a to 11a

The X component was obtained by blending each material of the X components shown in the following Table 1 to 5 at the ratios shown therein and mixing until homogeneous. Subsequently, the Y
10 component was added to the X component at the ratios shown in the said Tables to make the foam hair fixative. The proportion of amphoteric urethane resin to water-soluble resin designates dry weight (this applies to the Examples and Comparative Examples.)

The properties of foam hair fixatives thus obtained in the
15 Examples and Comparative Examples were evaluated in accordance with the following standards. The results are concurrently represented in the following Tables 1 to 5.

Touch

10 panelists conducted organoleptic tests on a sample of
20 black virgin hair (length:25cm, weight:5.0g) which had been coated with 0.8g of foam hair fixative and dried at normal room temperature for evaluation of touch as should inherently be endowed with as hair cosmetics. The evaluation standard is as follows.

: 9~10 persons who felt the sample of hair to be very soft to the touch.

: 6~8 persons who felt the sample of hair to be very soft to the touch.

5 : 2~5 persons who felt the sample of hair to be very soft to the touch.

×: 0~1 person who felt the sample of hair to be very soft to the touch.

Durability

10 Five curls, each consisting of black virgin hair (length:15cm, weight:3g) which had been coated with 0.6g of foam hair fixative, were prepared for each test piece, and held dried at a temperature of 50°C throughout the night. Then, the dried samples of hair were hung on a board marked with a graded scale in the thermo-hygrostat where the
15 temperature was maintained at 35°C and the moisture at 90%RH. The initial length of curl (a) and the length of curl (b) after five hours were measured to find the curl retentions in accordance with the following expression, wherein L is the length of a sample of hair which has been fully stretched.

20 Curl retention (%) $\{ (L - b) / (L - a) \} \times 100$

It is shown that as the values of the said curl retention approach 100%, the maintenance of curls is stronger and the durability more excellent. The evaluation standard is as follows.

90% or over

70% to under 90%

50% to under 70%

× Under 50%

Hair washability

5 A well-dressed sample of hair was prepared using black
virgin hair (length:15cm, weight:3g) which had been coated with 0.6g
of the foam hair fixative and dried at room temperature. Then, the said
sample of hair was slightly loosen using warm water at 40C for 30
seconds, then 0.4g of 10% shampoo solvent was applied to the
10 sample of hair which was then washed for 30 seconds. Then, it was
rinsed with warm water at 40C again to remove the shampoo
solvent, and dried sufficiently at 50C. Referring to the sample of hair
thus arranged, 10 panelists conducted organoleptic tests for hair-
setting ability and evaluated its hair washability which the foam hair
15 fixative should inherently be endowed with as hair cosmetics. The
evaluation standard is as follows.

: 9~10 persons who felt the sample of hair after drying to be
nil in setting ability but very excellent in hair washability.

: 6~8 persons who felt the sample of hair after drying to be
20 nil in setting ability but very excellent in hair washability.

: 2~5 persons who felt the sample of hair after drying to be
nil in setting ability but very excellent in hair washability.

×: 0~1 person who felt the sample of hair after drying to be nil
in setting ability but very excellent in hair washability.

Feel

For a sample of black virgin hair (length:25cm, weight:5.0g) which had been coated with 0.8g of the foam hair fixative and dried at the room temperature, 10 panelists conducted organoleptic tests and
5 evaluated the feel which the foam hair fixative should be inherently endowed with as hair cosmetics. The evaluation standard is as follows.

: 9~10 persons who felt the sample of hair after drying to have a very silky feel.

10 : 6~8 persons who felt the sample of hair after drying to have a very silky feel.

: 2~5 persons who felt the sample of hair after drying to have a very silky feel.

15 ×: 0~1 person who felt the sample of hair after drying to have a very silky feel.

20

Table 1
(Foam hair fixative) (Part)

	Example							
	1a	2a	3a	4a	5a	6a	7a	8a
Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
(Kind)	(a)	(a)	(a)	(a)	(a)	(a)	(a)	(a)
Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
De-ionized water	77.7	77.7	77.7	77.7	77.7	77.7	77.7	77.7
2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
Polyoxyethylene stearyl ether	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0
Coconut oil aliphatic acid diethanolamide	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8
Component Y	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0
Propellant (LPG)								
Touch								
Durability								
Hair washability								
Feel								

Table 2
(Foam Hair Fixative) (Part)

	Example									
	9a	10a	11a	12a	13a	14a	15a	16a		
Component X	Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
	(Kind)	(b)	(b)	(b)	(b)	(b)	(b)	(b)		
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	
	De-ionized water	77.7	77.7	77.7	77.7	77.7	77.7	77.7		
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-	
	Polyoxyethylene stearyl ether	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	
	Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0	
	Coconut oil aliphatic acid diethanolamide	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8	
	Component Y	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0	
Propellant (LPG)										
Touch										
Durability										
Hair washability										
Feel										

Table 3
(Foam hair fixative) (Part)

	Example									
	17a	18a	19a	20a	21a	22a	23a	24a		
Component X	Amphoteric urethane Resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
	(Kind)	(c)	(c)	(c)	(c)	(c)	(c)	(c)		
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)		
	De-ionized water	77.7	77.7	77.7	77.7	77.7	77.7	77.7		
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	Proper quantity	-		
	Polyoxyethylene stearyl ether	0.5	0.5	0.5	0.5	0.5	0.5	0.5		
	Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0		
	Coconut oil aliphatic acid diethanolamide	0.8	0.8	0.8	0.8	0.8	0.8	0.8		
	Component Y	8.0	8.0	8.0	8.0	8.0	8.0	8.0		
Propellant (LPG)										
Touch										
Durability										
Hair washability										
Feel										

Table 4

(Foam hair fixative) (Part)

	Comparative Example									
	1a	2a	3a	4a	5a	6a	7a	8a		
Component X	Amphoteric urethane resin	-	-	-	-	-	-	-	-	-
	(Kind)	-	-	-	-	-	-	-	-	-
	Water-soluble resin	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(8)
	De-ionized water	77.7	77.7	77.7	77.7	77.7	77.7	77.7	77.7	77.7
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-	-
	Polyoxyethylene stearyl ether	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
	Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0
	Coconut oil aliphatic acid diethanolamide	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8
	Component Y	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0	8.0
Propellant (LPG)										
Touch										
Durability										
Hair washability										
Feel										

Table 5
(Foam Hair Fixative) (Part)

	Comparative Examples		
	9a	10a	11a
Amphoteric urethane resin	3.0	3.0	3.0
(Types)	(a)	(b)	(c)
Water-soluble resin	-	-	-
(Kind)	-	-	-
Deionized water	77.7	77.7	77.7
2-amino-2-methyl-1-propanol	-	-	-
Polyoxyethylene stearyl ether	0.5	0.5	0.5
Ethanol	10.0	10.0	10.0
Coconut oil fatty acid diethanolamide	0.8	0.8	0.8
Component Y	8.0	8.0	8.0
Propellant (LPG)			
Touch			
Durability			
Hair washability			
Feel			

It is clear from the above Table 1 to 5 that the Examples of the foam hair fixative have very good touch, good feel, and ensure excellent hair washability because of the amphoteric urethane resin content, and are high in curl retention and excellent in durability
5 because of the water-soluble resin content.

Particularly, the Examples of the foam hair fixative, using anionic and amphoteric water-soluble resin, are extremely high in curl retention and excellent in durability. Also, the Examples of the foam hair fixative, using amphoteric urethane resin (b) structurally having
10 ethylene oxide chain(s), ensure excellent hair washability. Furthermore, the Examples of the foam hair fixative, using amphoteric urethane resin (c) structurally having polysiloxane bond(s), have very good feel.

In contrast, it is clear that the Comparative Examples 1a to 8a
15 of the foam hair fixative containing water-soluble resin are high in curl retention and excellent in durability and that the said examples containing no amphoteric urethane resin have very bad touch. Also, the Comparative Example 9a to 11a containing amphoteric urethane resin have very good touch and that the said examples containing no
20 water-soluble resin are low in curl retention and poor in durability.

Hair cosmetics (aerosol spray-type hair fixative)

Examples 1b to 24b, Comparative Examples 1b to 11b

The X component was obtained by blending each material of the X component shown in the following Table 6 to 10 at the ratios

shown therein and mixing until becoming homogeneous. Subsequently, the Y component was added to the X component at the ratios shown in the said Tables to make the aerosol spray-type hair fixative.

5 The properties of the aerosol spray type hair fixatives thus obtained in the Examples and Comparative Examples were evaluated in accordance with the standard for the aforementioned cosmetics for hair. The results are concurrently represented in the following Tables 6 to 10.

10

15

20

Table 6

(Aerosol spray-type hair fixative)		(Part)							
		Example							
		1b	2b	3b	4b	5b	6b	7b	8b
Component X	Amphoteric urethane Resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	(Kind)	(a)	(a)	(a)	(a)	(a)	(a)	(a)	(a)
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
	De-ionized water	7.0	7.0	7.0	7.0	7.0	7.0	7.0	7.0
	Diocetyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
	Ethanol	49.7	49.7	49.7	49.7	49.7	49.7	49.7	49.7
	Component Y	40.0	40.0	40.0	40.0	40.0	40.0	40.0	40.0
	Propellant (LPG)								
Touch									
Durability									
Hair washability									
Feel									

Table 7
(Aerosol spray-type hair fixative)
(Part)

	Examples									
	9b	10b	11b	12b	13b	14b	15b	16b		
Component X	Amphoteric urethane Resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
	(Kind)	(b)	(b)	(b)	(b)	(b)	(b)	(b)		
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-	
	De-ionized water	7.0	7.0	7.0	7.0	7.0	7.0	7.0	7.0	
	Dioctyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	
	Ethanol	49.7	49.7	49.7	49.7	49.7	49.7	49.7	49.7	
	Component Y	40.0	40.0	40.0	40.0	40.0	40.0	40.0	40.0	
	Propellant (LPG)									
Touch										
Durability										
Hair washability										
Feel										

Table 8
(Aerosol spray-type hair fixative)
(Part)

		Examples											
		17b	18b	19b	20b	21b	22b	23b	24b				
Component X	Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5				
	(Kind)	(c)	(c)	(c)	(c)	(c)	(c)	(c)	(c)				
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5				
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)				
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-				
	De-ionized water	7.0	7.0	7.0	7.0	7.0	7.0	7.0	7.0				
	Dioctyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3				
	Ethanol	49.7	49.7	49.7	49.7	49.7	49.7	49.7	49.7				
	Component Y	40.0	40.0	40.0	40.0	40.0	40.0	40.0	40.0				
	Propellant (LPG)												
Touch													
Durability													
Hair washability													
Feel													

Table 9
(Aerosol spray-type hair fixative) (Part)

		Comparative Example							
		1b	2b	3b	4b	5b	6b	7b	8b
Component X	Ampholeric urethane resin	-	-	-	-	-	-	-	-
	(Kind)	-	-	-	-	-	-	-	-
	Water-soluble resin	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
	De-ionized water	7.0	7.0	7.0	7.0	7.0	7.0	7.0	7.0
	Dioctyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
	Ethanol	49.7	49.7	49.7	49.7	49.7	49.7	49.7	49.7
Component Y		40.0	40.0	40.0	40.0	40.0	40.0	40.0	40.0
Propellant (LPG)									
Touch									
Durability									
Hair washability									
Feel									

Table 10
(Aerosol spray-type hair fixative) (Part)

	Comparative Example		
	9b	10b	11b
Component X	Amphoteric urethane resin	3.0	3.0
	(Kind)	(a)	(c)
	Water-soluble resin	-	-
	(Kind)	-	-
	2-amino-2-methyl-1-propanol	-	-
	De-ionized water	7.0	7.0
	Diocetyl sodium sulfosuccinate	0.3	0.3
	Ethanol	49.7	49.7
	Component Y	40.0	40.0
	Propellant (LPG)		
Touch			
Durability			
Hair washability			
Feel			

It is clear from the above Table 6 to 10 that the Examples of the aerosol spray-type hair fixative have very good touch, good feel, and ensure excellent hair washability because of containing amphoteric urethane resin, and are high in curl retention and excellent
5 in durability because of the water-soluble resin content.

In contrast, it is clear that the Comparative Examples 1b to 8b of the aerosol spray-type hair fixative containing no amphoteric urethane resin have very bad touch, and that the Comparative Examples 9b to 11b of the aerosol spray-type hair fixative containing
10 no water-soluble resin are low in curl retention and poor in durability.

Hair cosmetics (gel-like hair fixative)

Examples 1c to 24c, Comparative Examples 1c to 11c

The X component was obtained by blending each material of the X component shown in the following Table 11 to 15 at the ratios
15 shown therein and mixing until forming of viscous gel. Subsequently, the Y component obtained by blending each material at the ratios shown in the said Tables, was added to the above-mentioned X component and mixed until becoming homogeneous to make the gel type hair fixative.

20 The properties of the gel type hair fixatives thus obtained in the Examples and Comparative Examples were evaluated in accordance with the standard for the aforementioned cosmetics for hair. The results are concurrently represented in the following Tables 11 to 15.

Table 11

(Gel-like hair fixative) (Part)

	Examples							
	1c	2c	3c	4c	5c	6c	7c	8c
Component X	Thickening agent	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	Triethanolamine	1.1	1.1	1.1	1.1	1.1	1.1	1.1
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0
	De-ionized water	50.0	50.0	50.0	50.0	50.0	50.0	50.0
	Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5
Component Y	(Kind)	(a)	(a)	(a)	(a)	(a)	(a)	(a)
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	Proper quantity	-
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0
Touch	De-ionized water	34.4	34.4	34.4	34.4	34.4	34.4	34.4
Durability								
Hair washability								
Feel								

Table 12
(Gel-like hair fixative)
(Part)

		Examples							
		9c	10c	11c	12c	13c	14c	15c	16c
Component X	Thickening agent	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	Triethanolamine	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
	De-ionized water	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0
	Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
Component Y	(Kind)	(b)	(b)	(b)	(b)	(b)	(b)	(b)	(b)
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
Touch	De-ionized water	34.4	34.4	34.4	34.4	34.4	34.4	34.4	34.4
Durability									
Hair washability									
Feel									

Table 13

(Part)

(Gel-like hair fixative)

	Examples									
	17c	18c	19c	20c	21c	22c	23c	24c		
Component X	Thickening agent	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	Triethanolamine	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
	De-ionized water	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0
	Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
Component Y	(Kind)	(c)	(c)	(c)	(c)	(c)	(c)	(c)	(c)	(c)
	Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)	(8)
	2-amino-2-methyl-1-propanol	.	.	Proper quantity	.	.	Proper quantity	.	.	.
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
	De-ionized water	34.4	34.4	34.4	34.4	34.4	34.4	34.4	34.4	34.4
	Touch									
	Durability									
	Hair washability									
	Feel									

Table 14

(Gel-like hair fixative) (Part)

		Comparative Examples							
		1c	2c	3c	4c	5c	6c	7c	8c
		1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
Component X	Thickening agent	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1
	Triethanolamine	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
	Ethanol	50.0	50.0	50.0	50.0	50.0	50.0	50.0	50.0
	De-ionized water	-	-	-	-	-	-	-	-
Component Y	Amphoteric urethane resin	-	-	-	-	-	-	-	-
	(Kind)	-	-	-	-	-	-	-	-
	Water-soluble resin	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
	(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
	2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
	Ethanol	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
	De-ionized water	34.4	34.4	34.4	34.4	34.4	34.4	34.4	34.4
	Touch								
Durability									
Hair washability									
Feel									

Table 15
(Gel-like hair fixative) (Part)

	Component X	Comparative Examples		
		9c	10c	11c
	Thickening agent	1.5	1.5	1.5
	Triethanolamine	1.1	1.1	1.1
	Ethanol	5.0	5.0	5.0
	Deionized water	50.0	50.0	50.0
Component Y	Amphoteric urethane resin (Kind)	3.0	3.0	3.0
	Water-soluble resin (Kind)	(a)	(b)	(c)
		—	—	—
	2-amino-2-methyl-1-propanol	—	—	—
	Ethanol	5.0	5.0	5.0
	Deionized water	34.4	34.4	34.4
	Touch			
	Durability			
	Hair washability			
	Feel			

It is clear from the above Table 11 to 15 that the Examples of the gel-like hair fixative have very good touch, good feel, and ensure excellent hair washability because of containing amphoteric urethane resin, and are high in curl retention and excellent in durability because
5 of the water-soluble resin content.

In contrast, it is clear that the Comparative Examples 1c to 8c of the gel type hair fixative containing no amphoteric urethane resin have very bad touch, and that the Comparative Examples 9c to 11c of the gel type hair fixative containing no water-soluble resin are low in
10 curl retention and poor in durability.

Hair cosmetics (pump spray hair fixative)

Examples 1d to 24d, Comparative Examples 1d to 11d

Each component shown in the following Table 16 to 20 was blended at the ratios shown therein and was mixed until becoming
15 homogeneous to make the pump spray hair fixative.

The properties of the pump spray hair fixatives thus obtained in the Examples and Comparative Examples were evaluated in accordance with the standard for the aforementioned cosmetics for hair. The results are concurrently represented in the following Tables
20 16 to 20.

Table 16
(Pumping spray-type hair fixative) (Part)

	Examples							
	1d	2d	3d	4d	5d	6d	7d	8d
Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
(Kind)	(a)	(a)	(a)	(a)	(a)	(a)	(a)	(a)
Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5
(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
Diethyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0
De-ionized water	86.7	86.7	86.7	86.7	86.7	86.7	86.7	86.7
Touch								
Durability								
Hair washability								
Feel								

Table 17
(Pumping spray-type hair fixative) (Part)

	Examples									
	9d	10d	11d	12d	13d	14d	15d	16d		
Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
(Kind)	(b)	(b)	(b)	(b)	(b)	(b)	(b)	(b)		
Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)		
2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-		
Diethyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3		
Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0		
De-ionized water	86.7	86.7	86.7	86.7	86.7	86.7	86.7	86.7		
Touch										
Durability										
Hair washability										
Feel										

Table 18

(Pumping spray-type hair fixative)

(Part)

	Examples									
	17d	18d	19d	20d	21d	22d	23d	24d		
Amphoteric urethane resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
(Kind)	(c)	(c)	(c)	(c)	(c)	(c)	(c)	(c)		
Water-soluble resin	1.5	1.5	1.5	1.5	1.5	1.5	1.5	1.5		
(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)		
2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-		
Dioctyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3		
Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0		
De-ionized water	86.7	86.7	86.7	86.7	86.7	86.7	86.7	86.7		
Touch										
Durability										
Hair washability										
Feel										

Table 19
(Part)

(Pumping spray-type hair fixative)

	Comparative Examples							
	1d	2d	3d	4d	5d	6d	7d	8d
Amphoteric urethane resin	-	-	-	-	-	-	-	-
(Kind)	-	-	-	-	-	-	-	-
Water-soluble resin	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
(Kind)	(1)	(2)	(3)	(4)	(5)	(6)	(7)	(8)
2-amino-2-methyl-1-propanol	-	-	Proper quantity	-	-	-	Proper quantity	-
Dioctyl sodium sulfosuccinate	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Ethanol	10.0	10.0	10.0	10.0	10.0	10.0	10.0	10.0
De-ionized water	86.7	86.7	86.7	86.7	86.7	86.7	86.7	86.7
Touch								
Durability								
Hair washability								
Feel								

Table 20
(Pumping spray-type hair fixative) (Part)

	Comparative Examples		
	9d	10d	11d
Amphoteric urethane resin	3.0	3.0	3.0
(Kind)	(a)	(b)	(c)
Water-soluble resin	-	-	-
(Kind)	-	-	-
2-amino-2-methyl-1-propanol	-	-	-
Diethyl sodium sulfosuccinate	0.3	0.3	0.3
Ethanol	10.0	10.0	10.0
De-ionized water	86.7	86.7	86.7
Touch			
Durability			
Hair washability			
Feel			

It is clear from the above Table 16 to 20 that the Examples of the pump spray hair fixative have very good touch, good feel, and ensure excellent hair washability because of containing amphoteric urethane resin, and are high in curl retention and excellent in durability
5 because of containing water-soluble resin.

In contrast, it is clear that the Comparative Examples 1d to 8d of the pump spray hair fixative containing no amphoteric urethane resin have very bad touch, and that the Comparative Examples 9d to 11d of the pump spray hair fixative containing no water-soluble resin
10 are low in curl retention and poor in durability.

Skin care cosmetics (skin care lotion)

Examples 1e to 4e, Comparative Examples 1e to 5e

The X component was obtained by admixing each material of the X component shown in Table 21 and Table 22 mentioned later at the ratios shown therein and heating to 80°C. Also, the Y component
15 was obtained by admixing each material of the Y components shown in the said Tables at the ratios shown therein and heating to 80°C. Subsequently, the above-mentioned X and Y components were mixed and agitated at 80 °C for 30 minutes, and then a thickening agent was
20 added to the mixture and agitated until becoming homogeneous. After that, the (XY) mixture was cooled down to 40°C to make the skin care lotion.

The properties of the skin care lotion thus obtained in the Examples and Comparative Examples were evaluated in accordance

with the following standards. The results are concurrently represented in the following Tables 21 to 22.

Touch

10 panelists conducted practical application tests for
5 evaluation of the touch that the test pieces should be inherently endowed with as skin care cosmetics. The evaluation standard is as follows:

: 9 ~ 10 persons whose skin felt smooth and soft without feeling tight.

10 : 6 ~ 8 persons whose skin felt smooth and soft without feeling tight.

: 2 ~ 5 persons whose skin felt smooth and soft without feeling tight.

15 ×: 0 ~ 1 person whose skin felt smooth and soft without feeling tight.

Durability

10 panelists conducted practical application tests for
evaluation of the durability that the test pieces should be inherently endowed with as skin care cosmetics. The evaluation standard is as
20 follows:

: 9 ~ 10 persons who felt the applied cosmetics very lasting after lapse of 6 hours.

: 6 ~ 8 persons who felt the applied cosmetics very lasting after lapse of 6 hours.

: 2 ~ 5 persons who felt the applied cosmetics very lasting
after lapse of 6 hours.

×: 0 ~ 1 person who felt the applied cosmetics very lasting
after lapse of 6 hours.

5

10

15

20

(Skin care lotion)

Table 21

(Part)

	Examples			
	1e	2e	3e	4e
Component X	Octyl methoxycinnamate	7.5	7.5	7.5
	Ether polyoxystearate	1.0	1.0	1.0
	Emulsifiable glyceryl stearate	1.0	1.0	1.0
	Stearic acid	1.5	1.5	1.5
	Mixture of titanium dioxide and benzoic C ₁₂₋₁₅ alkyl	1.7	1.7	1.7
	Polyoxyethylene-additive dimethicone	0.5	0.5	0.5
Component Y	Amphoteric urethane resin	0.5	0.5	0.5
	(Kind)	(a)	(a)	(a)
	Water-soluble resin	0.5	0.5	0.5
	(Kind)	(1)	(5)	(7)
	De-ionized water	61.8	61.8	61.8
	Triethanolamine (99%)	4.0	4.0	4.0
	Antiseptic agent	Proper quantity	Proper quantity	Proper quantity
	Thickening agent (2%)	20.0	20.0	20.0
	2-amino-2-methyl-1-propanol	-	Proper quantity	Proper quantity
	Touch			
Durability				

Table 22
(Part)

(Skin care lotion)		Comparative Examples				
		1e	2e	3e	4e	5e
Component X	Octyl methoxycinnamate	7.5	7.5	7.5	7.5	7.5
	Polyoxystearate ether	1.0	1.0	1.0	1.0	1.0
	Emulsifiable glyceryl stearate	1.0	1.0	1.0	1.0	1.0
	Stearic acid	1.5	1.5	1.5	1.5	1.5
	Mixture of titanium dioxide and C ₁₂₋₁₅ alkyl benzoate	1.7	1.7	1.7	1.7	1.7
	Polyoxyethylene-additive Dimethicone	0.5	0.5	0.5	0.5	0.5
Component Y	Amphoteric urethane resin	-	-	-	-	1.0
	(Kind)	-	-	-	-	(a)
	Water-soluble resin	1.0	1.0	1.0	1.0	-
	(Kind)	(1)	(3)	(5)	(7)	-
	De-ionized water	61.8	61.8	61.8	61.8	61.8
	Triethanolamine (99%)	4.0	4.0	4.0	4.0	4.0
	Antiseptic agent	Proper quantity	Proper quantity	Proper quantity	Proper quantity	Proper quantity
	Thickening agent (2%)	20.0	20.0	20.0	20.0	20.0
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity	-
	Touch					
Durability						

It is clear from the above Table 21 and Table 22 that the Examples of the skin care lotion have very good touch because of the amphoteric urethane resin content and are excellent in durability
5 because of the water-soluble resin content.

In contrast, the Comparative Examples 1e to 4e of the skin care lotion containing no amphoteric urethane resin have very bad touch, and the compared example 5e of the skin care lotion containing no water-soluble resin is poor in durability.

10 Skin care cosmetics] (shaving cream)

Examples 1f to 4f, Comparative Examples 1f to 5f

The X component was obtained by admixing each material of the X components shown in the following Table 23 and Table 24 at the ratios shown therein and heating to 80°C. Also, the Y component was
15 obtained by admixing each material of the Y components at the ratios shown in the said Tables and heating to 80°C. Subsequently, the above-mentioned X and Y components were mixed at 80°C and cooled down to 40°C, and then proper amounts of antiseptic agent and perfume were added to the mixture to make the shaving cream
20 agent.

The properties of the shaving cream agent thus obtained in the Examples and Comparative Examples were evaluated in accordance with the standard for the aforementioned skin care cosmetics. The results are concurrently represented in the following
25 Tables 23 and Table 24.

Table 23

(Shaving cream agent) (Part)

	Examples			
	1f	2f	3f	4f
Component X	Stearic acid	8.0	8.0	8.0
	Mineral oil	2.0	2.0	2.0
	Isopropyl myristate	2.0	2.0	2.0
	Glycerol stearate	0.5	0.5	0.5
	Amphoteric urethane resin	0.3	0.3	0.3
Component Y	(Kind)	(a)	(a)	(a)
	Water-soluble resin	0.3	0.3	0.3
	(Kind)	(1)	(3)	(5)
	De-ionized water	72.7	72.7	72.7
	Thickening agent (2%)	10.0	10.0	10.0
Touch	Triethanolamine (99%)	4.2	4.2	4.2
	2-amino-2-methyl-1-propanol	-	Proper quantity	Proper quantity
Durability				

Table 24
(Shaving cream agent) (Part)

	Comparative Examples				
	1f	2f	3f	4f	5f
Component X	Stearic acid	8.0	8.0	8.0	8.0
	Mineral oil	2.0	2.0	2.0	2.0
	Isopropyl myristate	2.0	2.0	2.0	2.0
	Glycerol stearate	0.5	0.5	0.5	0.5
	Amphoteric urethane resin	-	-	-	0.6
Component Y	(Kind)	-	-	-	(a)
	Water-soluble resin	0.6	0.6	0.6	-
	(Kind)	(1)	(3)	(5)	(7)
	De-ionized water	72.7	72.7	72.7	72.7
	Thickening agent (2%)	10.0	10.0	10.0	10.0
Touch	Triethanolamine (99%)	4.2	4.2	4.2	4.2
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity
Durability					

It is clear from the above Table 23 and Table 24 that the Examples of the shaving cream agent have very good touch because of the amphoteric urethane resin content, and are excellent in durability because of the water-soluble resin content.

- 5 In contrast, the Comparative Examples 1f to 4f of the shaving cream agent containing no amphoteric urethane resin have very bad touch, and the Compared Example 5f of the shaving cream agent containing no water-soluble resin is poor in durability.

Skin care cosmetics (sunscreen lotion)

10 Examples 1g to 4g, Comparative Examples 1g to 5g

- The X component was obtained by admixing each material of the X components shown in the following Table 25 and Table 26 at the ratios shown therein and heating to 80°C. Also, the Y component was obtained by admixing each material of the Y components at the ratios
15 shown in the said Tables and heating to 80°C. Subsequently, the above-mentioned X and Y components were mixed at 80°C to make the sunscreen lotion.

- The properties of the sunscreen lotion thus obtained in the Examples and Comparative Examples were evaluated in accordance
20 with the standard for the aforementioned skin care cosmetics. The results are concurrently represented in the following Tables 25 and Table 26.

Table 25
(Sunscreen lotion)
(Part)

	Examples				
		1g	2g	3g	4g
Component X	Octyl methoxycinnamate	7.5	7.5	7.5	7.5
	Octyl palmitate	5.0	5.0	5.0	5.0
	Cetyl alcohol	1.0	1.0	1.0	1.0
	Polyethylene glycol monostearate	1.5	1.5	1.5	1.5
	Poly (oxyethylene oxypropylene) methylpolysiloxane copolymer	1.0	1.0	1.0	1.0
	Dimethylstearylamine	2.0	2.0	2.0	2.0
	Amphoteric urethane resin	1.0	1.0	1.0	1.0
Component Y	(Kind)	(a)	(a)	(a)	(a)
	Water-soluble resin	1.0	1.0	1.0	1.0
	(Kind)	(1)	(3)	(5)	(7)
	Purified water	69.0 5	69.0 5	69.0 5	69.0 5
	Triethanolamine (99%)	0.7	0.7	0.7	0.7
	Thickening agent (2%)	10.0	10.0	10.0	10.0
	Antiseptic agent	0.25	0.25	0.25	0.25
2-amino-2-methyl-1-propanol		-	Proper quantity	-	Proper quantity
Touch					
Durability					

Table 26
(Sunscreen lotion) (Part)

	Comparative Example				
	1g	2g	3g	4g	5g
Component X	Octyl methoxycinnamate	7.5	7.5	7.5	7.5
	Octyl palmitate	5.0	5.0	5.0	5.0
	Cetyl alcohol	1.0	1.0	1.0	1.0
	Polyethylene glycol monostearate	1.5	1.5	1.5	1.5
	Poly (oxyethylene oxypropylene) methylpolysiloxane copolymer	1.0	1.0	1.0	1.0
	Dimethylstearylamine	2.0	2.0	2.0	2.0
Component Y	Amphoteric urethane resin	-	-	-	2.0
	(Kind)	-	-	-	(a)
	Water-soluble resin	2.0	2.0	2.0	-
	(Kind)	(1)	(3)	(5)	(7)
	Purified water	69.0 5	69.0 5	69.0 5	69.0 5
	Triethanolamine (99%)	0.7	0.7	0.7	0.7
	Thickening agent (2%)	10.0	10.0	10.0	10.0
	Antiseptic agent	0.25	0.25	0.25	0.25
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity
	Touch				
Durability					

It is clear from the above Table 25 and Table 26 that the Examples of the sunscreen lotion have very good touch because of the amphoteric urethane resin content, and are excellent in durability because of the water-soluble resin content.

- 5 In contrast, the Comparative Examples 1g to 4g of the sunscreen lotion containing no amphoteric urethane resin have very bad touch, and that the compared example 5g of the sunscreen lotion containing no water-soluble resin is poor in durability.

Make-up cosmetics (nail polish)

10 Examples 1h to 4h, Comparative Example 1h to 5h

- As shown in Table 27 and Table 28 mentioned later, the pigment was dispersed in de-ionized water at the ratios shown in the said Tables, and then the other components were added at the ratios shown therein. After that, the solution was uniformly agitated, mixed,
15 and deaerated to make the nail polish.

The properties of the nail polish thus obtained in the Examples and Comparative Examples were evaluated in accordance with the following standards. The results are concurrently represented in the following Tables 27 and Table 28.

20 Touch

10 panelists conducted practical application tests for evaluation of the touch that the test pieces should be inherently endowed with as make-up cosmetics. The evaluation standard is as follows:

: 9 ~ 10 persons felt the applied area smooth and soft without feeling tight.

: 6 ~ 8 persons felt the applied area smooth and soft without feeling tight.

5 : 2 ~ 5 persons felt the applied area smooth and soft without feeling tight.

×: 0 ~ 1 person felt the applied area smooth and soft without feeling tight.

Durability

10 10 panelists conducted practical application tests for evaluation of the durability that the test pieces should be inherently endowed with as make-up cosmetics. The evaluation standard is as follows:

: 9 ~ 10 persons who felt the applied cosmetics very lasting after lapse of 6 hours.

: 6 ~ 8 persons who felt the applied cosmetics very lasting after lapse of 6 hours.

: 2 ~ 5 persons who felt the applied cosmetics very lasting after lapse of 6 hours.

20 ×: 0 ~ 1 person who felt the applied cosmetics very lasting after lapse of 6 hours.

Table 27
(Nail polish)
(Part)

		Examples			
		1h	2h	3h	4h
Aqueous phase	Amphoteric urethane resin	5.0	5.0	5.0	5.0
	(Kind)	(a)	(a)	(a)	(a)
	Water-soluble resin	5.0	5.0	5.0	5.0
	(Kind)	(1)	(3)	(5)	(7)
	De-ionized water	86.6	86.6	86.6	86.6
	Bentonite	0.6	0.6	0.6	0.6
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity
Pigment		2.5	2.5	2.5	2.5
Others	Perfume	0.1	0.1	0.1	0.1
	Antiseptic agent	0.1	0.1	0.1	0.1
	Silicone antifoamer	0.1	0.1	0.1	0.1
Touch					
Durability					

Table 28

(Nail polish) (Part)

		Comparative Examples				
		1h	2h	3h	4h	5h
Aqueous phase	Amphoteric urethane resin	-	-	-	-	10.0
	(Kind)	-	-	-	-	(a)
	Water-soluble resin	10.0	10.0	10.0	10.0	-
	(Kind)	(1)	(3)	(5)	(7)	-
	De-ionized water	86.6	86.6	86.6	86.6	86.6
	Bentonite	0.6	0.6	0.6	0.6	0.6
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity	-
Pigment		2.5	2.5	2.5	2.5	2.5
Others	Perfume	0.1	0.1	0.1	0.1	0.1
	Antiseptic agent	0.1	0.1	0.1	0.1	0.1
	Silicone antifoamer	0.1	0.1	0.1	0.1	0.1
Touch						
Durability						

It is clear from the above Table 27 and Table 28 that the Examples of the nail polish have very good touch because of the amphoteric urethane resin content, and are excellent in durability because of the water-soluble resin content.

- 5 In contrast, the Comparative Example 1h to 4h of the nail polish containing no amphoteric urethane resin have very bad touch, and that the Comparative Example 5h of the nail polish containing no water-soluble resin is poor in durability.

Make-up cosmetics (mascara)

- 10 Examples 1i to 4i, Comparative Examples 1i to 5i

- As shown in the following Table 29 and Table 30, propylene glycol, triethanolamine, thickening agent, antiseptic agent, and 2-amino-2-methyl-1-propanol were blended in refined water at the ratios shown in the said Tables and dissolved at 80°C, and then pigment
15 was dispersed therein to make an aqueous phase. Subsequently, each material of the Z component was blended at the ratios shown in the said Tables, and then dissolved at 80°C to make an oil phase. The oil phase was applied into the aqueous phase and emulsified by using a homogenizing mixer. After that, each material of the Y
20 component was slowly applied into the homogeneous emulsion at the ratios shown in the said Tables and agitated by using a homogenizing mixer, and then the emulsion was cooled down to normal temperature to make the mascara.

The properties of the mascara thus obtained in the Examples and Comparative Examples were evaluated in accordance with the standard for the aforementioned make-up cosmetics. The results are concurrently represented in the following Tables 29 and Table 30.

5

10

15

20

Table 29

(Mascara)

(Part)

	Examples	1i	2i	3i	4i
Component X	Propyleneglycol	5.0	5.0	5.0	5.0
	Triethanolamine	1.0	1.0	1.0	1.0
	Thickening agent (2%)	10.0	10.0	10.0	10.0
	Antiseptic agent	0.5	0.5	0.5	0.5
	Pigment	10.0	10.0	10.0	10.0
	Purified water	55.5	55.5	55.5	55.5
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity
Component Y	Amphoteric urethane resin	3.0	3.0	3.0	3.0
	(Kind)	(a)	(a)	(a)	(a)
	Water-soluble resin	3.0	3.0	3.0	3.0
	(Kind)	(1)	(3)	(5)	(7)
Component Z	Stearic acid	6.0	6.0	6.0	6.0
	Beeswax	6.0	6.0	6.0	6.0
Touch					
Durability					

Table 30
(Mascara) (Part)

		Comparative Examples				
		1i	2i	3i	4i	5i
Component X	Propyleneglycol	5.0	5.0	5.0	5.0	5.0
	Triethanolamine	1.0	1.0	1.0	1.0	1.0
	Thickening agent (2%)	10.0	10.0	10.0	10.0	10.0
	Antiseptic agent	0.5	0.5	0.5	0.5	0.5
	Pigment	10.0	10.0	10.0	10.0	10.0
	Purified water	55.5	55.5	55.5	55.5	55.5
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity	-
Component Y	Amphoteric urethane resin	-	-	-	-	6.0
	(Kind)	-	-	-	-	(a)
	Water-soluble resin	6.0	6.0	6.0	6.0	-
	(Kind)	(1)	(3)	(5)	(7)	-
	Stearic acid	6.0	6.0	6.0	6.0	6.0
Component Z	Beeswax	6.0	6.0	6.0	6.0	6.0
Touch						
Durability						

It is clear from the above Table 29 and Table 30 that the Examples of the mascara have very good touch because of the amphoteric urethane resin content, and are excellent in durability because of the water-soluble resin content.

- 5 In contrast, the Comparative Example 1i to 4i of the mascara containing no amphoteric urethane resin have very bad touch, and the Comparative Example 5i of the mascara containing no water-soluble resin is poor in durability.

Make-up cosmetics (foundation)

10 Examples 1j to 4j, Comparative Examples 1j to 5j

(1) Preparation of pigment

Each component shown in the following Table 31 and Table 32 was mixed at the ratios shown therein, and then milled through a grinder to prepare the pigment.

15 (2) Preparation of aqueous phase

After heating de-ionized water to 70°C, bentonite was added to the water for the purpose of swelling. Subsequently, sodium carboxymethyl cellulose previously dispersed in propylene glycol was added and dissolved therein. Subsequently, triethanol amine, paraoxy
20 methyl benzoate, 2-amino-2-methyl-1-propanol, and at least one of amphoteric urethane resin and water-soluble resin were added and dissolved therein to prepare the aqueous phase.

(3) Preparation of oil phase

Each component shown in the following Table 31 and Table 32 was mixed at the ratios shown therein, and then heated at 70°C to 80°C and dissolved to prepare the oil phase.

(4) Preparation of pigment dispersed solution

5 The aforementioned pigment was added to the above aqueous phase while agitating, and then treated through a colloid mill to prepare the pigment dispersed solution.

(5) Emulsification

 After heating the above pigment dispersed solution at 75°C
10 and the oil phase at 80°C respectively, the said oil phase was added to the said pigment dispersed solution while agitating, and then cooled to 45°C at which perfume was added, continuously agitated and cooled until reaching room temperature to prepare the foundation.

 The properties of the foundation thus obtained in the
15 Examples and Comparative Examples were evaluated in accordance with the standard for the aforementioned make-up cosmetics. The results are concurrently represented in the following Tables 31 and Table 32.

Table 31

(Foundation)

(Part)

		Examples			
		1j	2j	3j	4j
Oil phase	Stearic acid	2.4	2.4	2.4	2.4
	Propylene glycol monostearate	2.0	2.0	2.0	2.0
	Cetostearyl alcohol	0.2	0.2	0.2	0.2
	Liquid lanolin	2.0	2.0	2.0	2.0
	Liquid paraffin	3.0	3.0	3.0	3.0
	Isopropyl myristate	8.5	8.5	8.5	8.5
	Propyl paraoxybenzoate	Proper quantity	Proper quantity	Proper quantity	Proper quantity
Aqueous phase	Amphoteric urethane resin	0.3	0.3	0.3	0.3
	(Kind)	(a)	(a)	(a)	(a)
	Water-soluble resin	0.3	0.3	0.3	0.3
	(Kind)	(1)	(3)	(5)	(7)
	De-ionized water	63.5	63.5	63.5	63.5
	Carboxymethyl cellulose sodium	0.2	0.2	0.2	0.2
	Bentonite	0.5	0.5	0.5	0.5
	Propylene glycol	4.0	4.0	4.0	4.0
	Triethanolamine	1.1	1.1	1.1	1.1
	Methyl paraoxybenzoate	Proper quantity	Proper quantity	Proper quantity	Proper quantity
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity
Pigment	Titanium oxide	8.0	8.0	8.0	8.0
	Talc	4.0	4.0	4.0	4.0
	Coloring pigment	Proper Quantity	Proper Quantity	Proper Quantity	Proper Quantity
Perfume		Proper quantity	Proper quantity	Proper quantity	Proper quantity
Touch					
Durability					

(Foundation)

Table 32

(Part)

		Comparative Examples				
		1j	2j	3j	4j	5j
Oil phase	Stearic acid	2.4	2.4	2.4	2.4	2.4
	Propylene glycol monostearate	2.0	2.0	2.0	2.0	2.0
	Cetostearyl alcohol	0.2	0.2	0.2	0.2	0.2
	Liquid lanolin	2.0	2.0	2.0	2.0	2.0
	Liquid paraffin	3.0	3.0	3.0	3.0	3.0
	Isopropyl myristate	8.5	8.5	8.5	8.5	8.5
	Propyl paraoxybenzoate	Proper quantity	Proper quantity	Proper quantity	Proper quantity	Proper quantity
Aqueous phase	Amphoteric urethane resin	-	-	-	-	0.6
	(Kind)	-	-	-	-	(a)
	Water-soluble resin	0.6	0.6	0.6	0.6	-
	(Kind)	(1)	(3)	(5)	(7)	-
	De-ionized water	63.5	63.5	63.5	63.5	63.5
	Carboxymethyl cellulose sodium	0.2	0.2	0.2	0.2	0.2
	Bentonite	0.5	0.5	0.5	0.5	0.5
	Propyleneglycol	4.0	4.0	4.0	4.0	4.0
	Triethanolamine	1.1	1.1	1.1	1.1	1.1
	Methyl paraoxybenzoate	Proper quantity	Proper quantity	Proper quantity	Proper quantity	Proper quantity
	2-amino-2-methyl-1-propanol	-	Proper quantity	-	Proper quantity	-
Pig-ment	Titanium oxide	8.0	8.0	8.0	8.0	8.0
	Talc	4.0	4.0	4.0	4.0	4.0
	Coloring pigment	Proper Quantity	Proper Quantity	Proper Quantity	Proper Quantity	Proper Quantity
Perfume		Proper quantity	Proper quantity	Proper quantity	Proper quantity	Proper quantity
Touch						
Durability						

It is clear from the above Table 31 and Table 32 that the Examples of the foundation have very good touch because of the amphoteric urethane resin content, and are excellent in durability because of the water-soluble resin content.

5 In contrast, the Comparative Examples 1j to 4j of the foundation containing no amphoteric urethane resin have very bad touch, and the compared example 5j of the foundation containing no water-soluble resin is poor in durability.

Effects of the Invention

10 As described above, in the cosmetics of the present invention, using as a base resin amphoteric urethane resin containing carboxyl group(s) and tertiary amino group(s) in one molecule, originally antithetic physical properties such as setting ability and touch, combing ability and resistance to flaking are compatible with
15 each other by the elasticity and toughness that the urethane resin possesses. Furthermore, as against neutral water, the said cosmetics ensure excellent water resistance because of the ion bond(s) between carboxyl group(s) and tertiary amino group(s). On the other hand, as against surface-active agent solution such as shampoo, etc., the said
20 cosmetics ensure excellent hair washability with dissociation of the said ion bond(s), and at the same time, the above-mentioned cationic tertiary amino group(s) interacts on the surfaces of the negatively charged hair to ensure excellent adhesion. Moreover, the cosmetics of the present invention use both water-soluble resin and amphoteric

urethane resin, therefore the problem of the durability which is a weak point of amphoteric urethane resin can be solved by the use of water-soluble resin, and also the problem of the touch which is a weak point of water-soluble resin can be solved by the use of amphoteric urethane resin, and thereby the said cosmetics are provided with antithetic physical properties such as touch and durability which are required for cosmetics.

Additionally, the durability may be further improved by using nonionic resin, anionic resin, cationic resin or amphoteric resin as said water-soluble resin.

The introduction into the structure of the said amphoteric urethane resin of the structural unit which may be derived from ethylene oxide as nonionic hydrophilic constituent may provide sufficient hydrophilic nature, which ensures improvement of hair washability especially when it is used as hair cosmetic material.

Additionally, the introduction of polysiloxane bond(s) into the structure of the said amphoteric urethane resin can improve the feel which may be felt especially when it is used as hair cosmetic material.

WHAT IS CLAIMED IS:

1. Cosmetics characterized in that it contains amphoteric urethane resin having carboxyl group(s) and
5 tertiary amino group(s), and water-soluble resin in one molecule.
2. The cosmetics as defined in claim 1, wherein the water-soluble resin is nonionic resin.
- 10 3. The cosmetics as defined in claim 1, wherein the water-soluble resin is anionic resin.
4. The cosmetics as defined in claim 1, wherein the water-soluble resin is cationic resin.
- 15 5. The cosmetics as defined in claim 1, wherein the water-soluble resin is amphoteric resin.
6. The cosmetics as defined in any of claims 1 to 5, wherein the
20 amphoteric urethane resin has in its structure a structural unit that is derived from ethylene oxide.
7. The cosmetics as defined in any of claims 1 to 5, wherein the amphoteric urethane resin has in its structure polysiloxane bond(s).

8. The cosmetics as defined in any of claims 1 to 7, wherein the amphoteric resin is aqueous liquid.

- 5 9. The cosmetics as defined in any of claims 1 to 8, wherein it is at least one selected from the group consisting of hair cosmetics, skin care cosmetics and make-up cosmetics.

10

15

20

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 00/21872

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61K7/06 A61K7/48 A61K7/11 A61K7/15 A61K7/043
A61K7/02 A61K7/032

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, PAJ, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 775 483 A (OREAL) 28 May 1997 (1997-05-28) claims 1,6,8,18 ----	1-9
X	EP 0 793 957 A (OREAL) 10 September 1997 (1997-09-10) claims 1,6-11,20 ----	1-6,8,9
E	US 6 126 929 A (MOUGIN NATHALIE) 3 October 2000 (2000-10-03) claims 1,5-9,32-39 -----	1-5,8,9

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

20 November 2000

Date of mailing of the international search report

27/11/2000

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Beyss, E

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/US 00/21872

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0775483	A	28-05-1997	FR 2740336 A	30-04-1997
			FR 2740330 A	30-04-1997
			BR 9604455 A	23-06-1998
			CA 2188691 A	28-04-1997
			JP 9169615 A	30-06-1997
			JP 2000136117 A	16-05-2000
			US 5972354 A	26-10-1999
EP 0793957	A	10-09-1997	FR 2745494 A	05-09-1997
			AT 175109 T	15-01-1999
			BR 9707855 A	27-07-1999
			CA 2199053 A	04-09-1997
			CN 1212617 A	31-03-1999
			DE 69700082 D	11-02-1999
			DE 69700082 T	27-05-1999
			ES 2129998 T	16-06-1999
			WO 9732566 A	12-09-1997
			JP 11507396 T	29-06-1999
			PL 328600 A	01-02-1999
			US 6010686 A	04-01-2000
US 6126929	A	03-10-2000	CA 2249478 A	23-07-1998
			EP 0909157 A	21-04-1999
			WO 9831329 A	23-07-1998